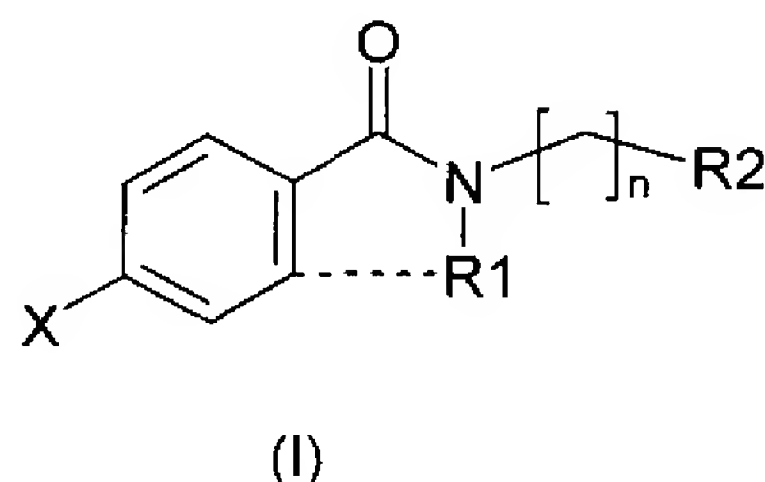


AMENDMENTS TO THE CLAIMS:

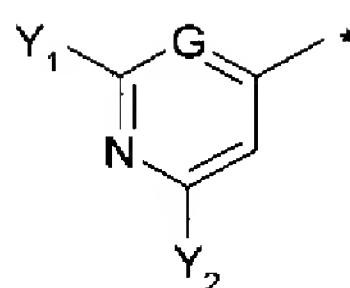
Please amend the claims as follows:

1. (Currently amended) A compound of Formula (I) or a salt, or solvate, ~~or physiologically functional derivative thereof~~:



wherein:

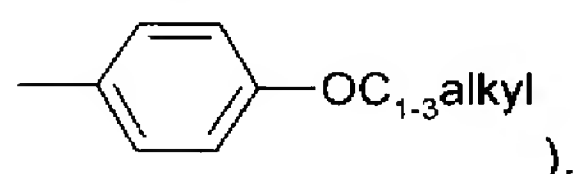
- R1 is hydrogen or C₁₋₆alkyl or as indicated by the dotted line is fused to the phenyl forming a 5 or 6 membered ring, optionally containing a double bond;
- n is 0, 1, 2, 3 or 4;
- R2 is aryl, optionally substituted by one or two groups selected from the group consisting of halogen, NH₂, hydroxy, cyano, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, aryl, aryloxy, C₁₋₄alkoxycarbonyl, C₁₋₄alkylsulfonyl and a group R₃R₄NSO₂ (wherein R₃ and R₄ are independently hydrogen or C₁₋₄alkyl), (CH₂)₀₋₃NHCOOC₁₋₄alkyl, and a 5- or 6-membered heteroaryl group;
- or n is 0 and R1 and R2, together with the nitrogen atom to which they are joined, form a 5- or 6-membered monocyclic heterocyclic ring or a 9- or 10-membered bicyclic heterocyclic ring wherein at least the ring which contains the nitrogen atom to which R1 and R2 are joined is non-aromatic, and wherein the 5- or 6-membered monocyclic heterocyclic ring or the 9- or 10-membered bicyclic heterocyclic ring is optionally substituted by one or two groups selected from the group consisting of halogen, hydroxy, cyano, oxo, C₁₋₄alkyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, aryl, aryloxy and C₁₋₄alkoxycarbonyl; and
- X is indazolyl, pyrazolyl or a group



wherein

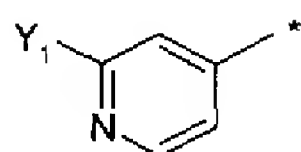
G is CH or N; and

Y₁ and Y₂ are independently hydrogen, halogen and a group NR₅R₆ (wherein R₅ and R₆ are independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl, or



2. (Original) A compound as claimed in claim 1, wherein R₁ is hydrogen.
3. (Currently amended) A compound as claimed in claim 1 ~~or claim 2~~, wherein n is 1 or 2.
4. (Currently amended) A compound as claimed in claim 1 ~~any of claims 1-3~~, wherein R₂ is aryl, optionally substituted by one or two groups selected from the group consisting of halogen and C₁₋₄alkoxy.
5. (Currently amended) A compound as claimed in claim 1 ~~or claim 2~~, wherein n is 0 and R₁ and R₂, together with the nitrogen atom to which they are joined, form a 6-membered monocyclic heterocyclic ring or a 10-membered bicyclic heterocyclic ring wherein at least the ring which each contains the nitrogen atom to which R₁ and R₂ are joined is non-aromatic, wherein the 6-membered monocyclic heterocyclic ring or 10-membered bicyclic heterocyclic ring are both optionally substituted by one or two groups selected from oxo, C₁₋₄alkyl, phenyl and C₁₋₄alkoxycarbonyl.
6. (Currently amended) A compound as claimed in claim 1 ~~any of claims 1-5~~, wherein X is indazolyl or pyrazolyl.

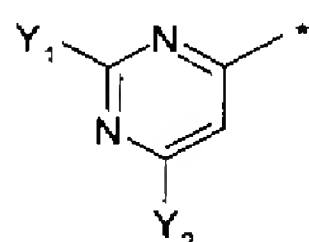
7. (Currently amended) A compound as claimed in claim 1 ~~any of claims~~



~~4-5~~, wherein X is a group:

wherein Y₁ is hydrogen or halogen.

8. (Currently amended) A compound as claimed in claim 1 ~~any of claims 1-5~~, wherein X is a group:



wherein one of Y₁ and Y₂ is hydrogen, and the other is hydrogen, halogen or a group NR₅R₆ wherein R₅ and R₆ are independently hydrogen, C₁₋₆alkyl or C₂₋₆alkenyl.

9. (Currently amended) A compound as claimed in claim 1, wherein said compound is selected from the group consisting of ~~which is:~~

N-benzyl-4-(4-pyridinyl)benzamide;

N-(2-phenylethyl)-4-(4-pyridinyl)benzamide;

N-(3-methoxybenzyl)-4-(4-pyridinyl)benzamide;

N-(3-methoxybenzyl)-4-(1H-pyrazol-4-yl)benzamide;

4-(2-chloro-4-pyridinyl)-N-(3-methoxybenzyl)benzamide;

4-(2-amino-4-pyrimidinyl)-N-(3-methoxybenzyl)benzamide;

N-(3-methoxybenzyl)-4-(4-pyrimidinyl)benzamide;

4-[6-(allylamino)-4-pyrimidinyl]-N-(3-methoxybenzyl)benzamide;

4-[6-amino-4-pyrimidinyl]-N-(3-methoxybenzyl)benzamide;

4-(1H-indazol-5-yl)-N-(3-methoxybenzyl)benzamide;

4-(2-amino-4-pyrimidinyl)-N-[[3-(methoxy)phenyl]methyl]benzamide;

4-(2-[[4-(methoxy)phenyl]amino]-4-pyrimidinyl)-N-[[3-(methoxy)phenyl]methyl]benzamide;

4-(2-amino-4-pyrimidinyl)-N-[(2-chlorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(4-fluorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(4-chlorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(2-fluorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[[2-(methyloxy)phenyl]methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(2-methylphenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[[4-(methyloxy)phenyl]methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-(2,3-dihydro-1H-inden-1-yl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(1R)-1,2,3,4-tetrahydro-1-naphthalenyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[[3-(trifluoromethyl)phenyl]methyl]benzamide;
 1,1-dimethylethyl({3-[[[4-(2-amino-4-pyrimidinyl)phenyl]carbonyl]amino)methyl]phenyl)methyl}carbamate;
 4-(2-amino-4-pyrimidinyl)-N-[(3-bromophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(3-chlorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(3-fluorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-(phenylmethyl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(1S)-1-[4-(methyloxy)phenyl]ethyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(1S)-1-phenylpropyl]benzamide;
 6-(2-amino-4-pyrimidinyl)-2-(phenylmethyl)-1(2H)-isoquinolinone;
 4-(2-amino-4-pyrimidinyl)-N-[(3-hydroxyphenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-({3-[(difluoromethyl)oxy]phenyl}methyl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-(1-methyl-1-phenylethyl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(3,5-dichlorophenyl)methyl]benzamide;
 4-(2-amino-4-pyrimidinyl)-N-(4-biphenylmethyl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(1R)-1-phenylethyl]benzamide;
 1,1-dimethylethyl{3-[1-({[4-(2-amino-4-pyrimidinyl)phenyl]carbonyl}amino)ethyl]phenyl}carbamate;
 N-[(2-aminophenyl)methyl]-4-(2-amino-4-pyrimidinyl)benzamide;
 4-(2-amino-4-pyrimidinyl)-N-[(1S)-1,2,3,4-tetrahydro-1-naphthalenyl]benzamide;
 5-(2-amino-4-pyrimidinyl)-2-(phenylmethyl)-2,3-dihydro-1H-isoindol-1-one;
 4-(2-amino-4-pyrimidinyl)-N-[(1R)-1-[3-(methyloxy)phenyl]ethyl]benzamide; and
 4-(2-amino-4-pyrimidinyl)-N-[(1R)-1-phenylpropyl]benzamide;

or a salt, or solvate ~~or physiologically functional derivative~~ thereof.

10-11. (Cancelled).

12. (Currently amended) A method of treating a disorder in a mammal, said disorder being mediated by inappropriate ROCK-1 activity, comprising: administering to said mammal a therapeutically effective amount of a compound as defined in claim 1~~any of claims 1-8~~.

13. (Cancelled).

14. (Currently amended) A pharmaceutical composition comprising a therapeutically effective amount of a compound as defined in claim 1~~any of claims 1-8~~ and one or more of pharmaceutically acceptable carriers, diluents and excipients.